## WHAT IS CLAIMED IS:

I	1. A method for screening a library of monome	er domains or multimers		
2	comprising monomer domains for binding affinity to multiple ligands, the method comprising			
3	contacting a library of monomer domains or multimers of monomer domains			
4	4 to multiple ligands; and	to multiple ligands; and		
5	selecting monomer domains or multimers that bind	to at least one of the		
6	6 ligands.			
1	1 2. The method of claim 1, comprising			
2	•	multiple ligands;		
3				
4		<del>-</del>		
5	domains to form a library of multimers, each comprising a selected monomer domain and a			
6	second monomer domain;			
7	7 iv. contacting the library of multimers to the man	ultiple ligands to form a		
8	plurality of complexes, each complex comprising a multimer and a ligand; and			
9	v. selecting at least one complex.			
1	1 3. The method of claim 2, the method further of	comprising		
2	linking the multimers of the selected complexes to a	a library of monomer		
3	domains or multimers to form a second library of multimers, each	domains or multimers to form a second library of multimers, each comprising a selected		
4	multimer and at least a third monomer domain;			
5	contacting the second library of multimers to the m	ultiple ligands to form a		
6	plurality of second complexes; and			
7	selecting at least one second complex.			
1	1 4. The method of claim 2, wherein the identity	of the ligand and the		
2	2 multimer is determined.			
1	1 5. The method of claim 1, wherein a library of	monomer domains is		
2	,	monomial domining is		
l	6. The method of claim 1, wherein a library of	multimers is contacted to		
2	2 multiple ligands.			

1	7.	The method of claim 1, wherein the multiple ligands are in a mixture.	
1	8.	The method of claim 1, wherein the multiple ligands are in an array.	
1	9.	The method of claim 1, wherein the multiple ligands are in or on a cell	
2	or tissue.		
1	10.	The method of claim 1, wherein the multiple ligands are immobilized	
2	on a solid support.		
1	11.	The method of claim 1, wherein the ligands are polypeptides.	
1	12.	The method of claim 12, wherein the polypeptides are expressed on the	
2	surface of phage.		
1	13.	The method of claim 1, wherein the monomer domain or multimer	
2	library is expressed on the surface of phage.		
1	14.	The method of claim 1, wherein the monomer domain is a LDL	
2	receptor type A mo	nomer domain.	
1	15.	The method of claim 1, wherein the monomer domain is an EGF	
2	monomer domain.		
1	16.	The method of claim 1, wherein the library of multimers is expressed	
2	on the surface of phage to form library-expressing phage and the ligands are expressed on the		
3	surface of phage to form ligand-expressing phage, and the method comprises		
4	conta	acting library-expressing phage to the ligand-expressing phage to form	
5		hage/library-expressing phage pairs;	
6	remo	oving ligand-expressing phage that do not bind to library-expressing or	
7	removing library-expressing phage that do not bind to ligand-expressing phage; and		
8		ting the ligand-expressing phage/library-expressing phage pairs.	
1	17.	The method of claim 16, further comprising isolating polynucleotides	
2	from the phage pair	s and amplifying the polynucleotides to produce a polynucleotide hybrid	
3		leotides from the ligand-expressing phage and the library-expressing	
4	phage.		

1	18. The method of claim 17, comprising isolating polynucleotide hybrids		
2	from a plurality of phage pairs, thereby forming a mixture of polynucleotide hybrids.		
1	19. The method of claim 18, comprising		
2	contacting the mixture of hybrid polynucleotides to a cDNA library under		
3	conditions to allow for polynucleotide hybridization, thereby hybridizing a hybrid		
4	polynucleotide to a cDNA in the cDNA library; and		
5	determining the nucleotide sequence of the hybridized hybrid polynucleotide		
6	thereby identifying a monomer domain that specifically binds to the polypeptide encoded by		
7	the cDNA.		
1	20. The method of claim 1, wherein the monomer domain library is		
2	expressed on the surface of phage to form library-expressing phage and the ligands are		
3	expressed on the surface of phage to form ligand-expressing phage, and the selected		
4	complexes comprise a library-expressing phage bound to a ligand-expressing phage and the		
5	method comprises:		
6	dividing the selected monomer domains or multimers into a first and a second		
7	portion,		
8	linking the monomer domains or multimers of the first portion to a solid		
9	surface and contacting a phage-displayed ligand library to the monomer domains or		
10	multimers of the first portion to identify target ligand phage that binds to a monomer domain		
11	or multimer of the first portion;		
12	infecting phage displaying the monomer domains or multimers of the second		
13	portion into bacteria to express the phage; and		
14	contacting the target ligand phage to the expressed phage to form phage pairs		
15	comprised of a target ligand phage and a phage displaying a monomer domain or multimer.		
1	21. The method of claim 20, further comprising isolating a polynucleotide		
2	from each phage of the phage pair, thereby identifying a multimer or monomer domain that		
3	binds to the ligand in the phage pair.		
1	22. The method of claim 23, further comprising amplifying the		
2	polynucleotides to produce a polynucleotide hybrid comprising polynucleotides from the		
3	target ligand phage and the library phage.		

1	23. The method of claim 20, comprising isolating and amplifying		
2	polynucleotide hybrids from a plurality of phage pairs, thereby forming a mixture of		
3	polynucleotide hybrids.		
1	24. The method of claim 23, comprising		
2	contacting the mixture of hybrid polynucleotides to a cDNA library under		
3	conditions to allow for hybridization, thereby hybridizing a hybrid polynucleotide to a cDNA		
4	in the cDNA library; and		
5	determining the nucleotide sequence of the associated hybrid polynucleotide,		
6	thereby identifying a monomer domain that specifically binds to the ligand encoded by the		
7	cDNA associated cDNA.		